Roles of Mammalian Coro1B in Actin Dynamics and Cell Migration

Liang CAI, James E. Bear

Department of Cell and Developmental Biology, University of North Carolina at Chapel Hill, Chapel Hill, NC, USA

BACKGROUND

Cell migration is a highly integrated multi-step process that orchestrates embryonic morphogenesis; contributes to tissue repair and regeneration; takes part in immune response and neuron development; and drives disease progression in cancer, mental retardation, atherosclerosis, and arthritis. Though cell motility has been studied for decades, and molecular basis of cell migration has progressed rapidly over the recently few years, there are still many unresolved issues regarding how cell maintain the polarity, how adhesions disperse, how cells recognize their targets, how components are integrated temporally and spatially across the cell. (*Science*, 302:1704.) Cell migration requires the participation of a dynamic and interactive actin cytoskeleton that acts to deform cellular membranes. Many factors have been reported to be concerned with these 'actin-based motility processes'.

Coronins are a conserved family of F-actin binding proteins that contain WD40 repeats. Yeast Saccharomyces cerevisiae and Dictyostelium discoideum each have a single gene encoding Coronin; but in mammals, this gene family has expanded to include seven genes: CORO1A, CORO1B, CORO1C, CORO1D, CORO2A, CORO2B and POD (based on the HGNC nomenclature).

Coro1B, first identified as a downstream target of PKC in 1999, is expressed in many tissue and cell lines. Some primary experiments show that Coro1B is localized to the region of dynamic actin at the leading edge and weakly along stress fibers, which indicates possible roles of Coro1B in actin dynamics and cell migration.

RESULTS AND FUTURE WORKS Fig.4 Antibody tests Fig.6 Cell state influences the location of hCoro1B-EGFP Suspended Cell Rat2: Rat2: hCoro1B-EGFP Rat2 thCoro1B-EGFP S P S P S P S RIPA buffer anti-hCoro1B-1 Fig.7 Diagram of Retrovirus constructs anti-hCoro1B-2 IB: anti-hPOD SET.I hCoro1B-mRFP IB: anti-GFP EGFP MT hCoro1B-Coiled-Coil IB: anti-p34 SET.II EGFP hCoro1B-EGFP Fig.5 Trial of hCoro1B IP GFP h1B-deltWD40-EGFF How hCoro1B binds to actin? EGFP WD40-EGFP SET.III IP: anti-GFP EGFP mCoro1B-EGFP (For siRNA knockdown experiments)

BIOINFORMATICS ANALYSIS

Tab.1 Sequence similarity based on DNA (upper)/amino acid(lower) data

h1A	m1A	r1A		h1B	m1B	r1B	
1	0.83	0.83	h1B	1	0.82	0.79	
0.94	1	0.96	m1B	0.94	1	0.91	
0.94	0.99	1	r1B	0.92	0.97	1	
h2A	m2A		h2B	m2B		h1C	m1C
1	0.51	h2B	1	0.68	h1C	1	0.69
0.85	1	m2B	0.98	1	m1C	0.97	1
h1D	m1D	r1D_		hPO	D mP(DD rl	POD
1	0.51	0.40	hPOD	1	0.6	4 (0.52
0.47	1	0.49	mPOD	0.86	1	(0.42
0.46	0.95	1	rPOD	0.37	0.4	0	1
	1 0.94 0.94 h2A 1 0.85 h1D 1 0.47	1 0.83 0.94 1 0.94 0.99 h2A m2A 1 0.51 0.85 1 h1D m1D 1 0.51 0.47 1	1	1	1 0.83 0.83 h1B 1 0.94 0.94 0.99 1 r1B 0.92 0.92 0.94 0.91 0.92 0.86 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 1 0.98 0.98 1 0.49 0.86	1	1

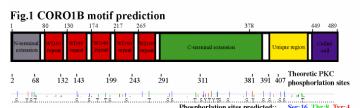


Fig.2 Phylogenetic tree of coronin family

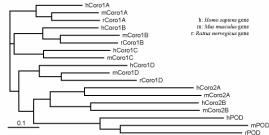


Fig.3 hCoro1B 3-D structure predicted by SWISS-MODEL

